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**ANALYSIS  
OF  
SCROTAL SWELLINGS  
IN  
GOVT. RAJAJI HOSPITAL  
MADURAI**



**THE TAMILNADU DR. M.G.R. MEDICAL  
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# CERTIFICATE

This is to certify that dissertation entitled “ANALYSIS OF SCROTAL SWELLINGS IN GOVT.RAJAJI HOSPITAL, MADURAI” Submitted by Dr. K. K. Saravanan to the Tamil Nadu Dr. M.G.R Medical University , Chennai, is in partial fulfillment of the requirement for the award of M.S degree Branch – I (General Surgery) and is a bonafide research work carried out by him under direct supervision and guidance.

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# DECLARATION

This is consolidated report on “**ANALYSIS OF SCROTAL SWELLINGS IN GOVT.RAJAJI HOSPITAL MADURAI**” based on 324 cases treated at Govt. Rajaji Hospital, Madurai, during the period June 2004 to September 2006.

This is submitted to the **Tamilnadu Dr. M.G.R. Medical University, Chennai** in partial fulfillment of the rules and regulations for the **M.S.** Degree Examination in General Surgery.

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# INTRODUCTION

The scrotum is a sac of skin and involuntary muscles enclosing the testis, epididymis, vas deferens and spermatic vessels is a phylogenetic sophistication of the genito urinary system. The scrotum functions to provide an apt environment to the testis for optimal spermatogenesis, which is so important for perpetuation of a race. In other words scrotum is the thermoregulator of spermatogenesis.

The importance of scrotum, its contents and their pathologies can be assumed from it being considered as the tenth compartment of the abdomen. The high incidence of male infertility and other morbidities associated with scrotal pathologies make a detailed study into them and their management worthwhile.

Most of the patients with disease of the scrotum and its contents present with swelling of the scrotum and little more few other symptoms and signs. This study intends to look into the various pathologies of the scrotum and its contents, which present as scrotal swelling; and to highlight upon the best way to approach them; in order to restore the anatomy and physiology to the maximum possible level.

## **AIMS OF STUDY**

1. To study the various causes for scrotal swellings and their pattern of incidence in and around Madurai.
2. To study the various clinical presentations of patients with swelling of the scrotum.
3. To analyse the diagnostic and therapeutic modalities adopted in the management of scrotal swellings in our hospital and their outcome.
4. To study the relationship of certain scrotal swelling with male infertility.
5. To highlight on the recent trends in the diagnosis and management of the various scrotal pathologies.

## MATERIALS AND METHODS

324 patients with scrotal swelling were admitted in the Department of General surgery, Government Rajaji-Hospital-Madurai Medical College during the period of June-04 to September-06.

Under the guidance of Professor **Dr. S. Vijayalakshmi M.S.**, Detailed clinical evaluation with necessary investigations to identify the cause and its effects were done.

The different methods of available treatment and their outcome were studied. In selected cases pre-operative and postoperative clinical photographs were taken, and subsequent follow up was done.



# **ANATOMY OF THE SCROTUM**

The scrotum is a pendulous bag of skin & fascia designed to lodge the testis. The scrotal wall is composed of

- a. Skin
- b. Subcutaneous fascia Dartos muscle
- c. External spermatic fascia
- d. Cremasteric fascia
- e. Internal spermatic fascia
- f. Tunica vaginalis

## **BLOOD SUPPLY OF SCROTUM**

The scrotum is supplied by

- a. Superficial and deep external pudendal branches of femoral artery.
- b. Medial and lateral scrotal branches of the internal pudendal artery.
- c. Cremasteric artery from inferior epigastric artery

## **NERVE SUPPLY OF SCROTUM**

Ventral part - Ilioinguinal nerve (L1)

Dorsal part - scrotal branches of pudendal nerve (S2,S3)

Perineal branches of posterior cutaneous nerve of thigh.

Cremaster - Genital branch of genitogemoral nerve (L1,L2)

## **LYMPHATIC DRAINAGE**

**Lymphatic vessels from the testis run upwards in the spermatic cord and alongside the veins end in the para aortic lymph nodes at the level of the first lumbar vertebra (Smith, 1983).**

Lymphatic drainage of the scrotum takes place to the inguinal glands.

### ***CONTENTS OF THE SCROTUM***

Scrotum contain both the testis and the corresponding spermatic cords.

### ***TESTIS***

Testis lies in the scrotum with its long axis oblique and upper pole tilted forward, with vas deferens and epididymis posteriorly. Size is about 5cm x 2.5cm x 3cm.

Testis is covered by bluish white “Tunica albuginea” which invades the interior to form the mediastinum along its posterior border.

150 to 200 septae divide the interior into lobules, “Rete Testis” from which vasa efferentia enter the head of the epididymis.

### ***EPIDIDYMIS***

It is coiled structure containing a single epididymal tubule and attached to the posterolateral surface of each testis. It has a head, body & tail. The canal of epididymis leaves the tail and continues as vas deferens.

## **SPERMATIC CORD**

**It extends from the deep inguinal ring to the posterosuperior border of the testicle, surrounded by three fibrous coats derived from the abdominal wall during the descent of the testis**

- a. External spermatic fascia**
- b. Cremasteric fascia**
- c. Internal spermatic fascia**

**along with peritoneum and preperitoneal fat contains vas deferens**

**Spermatic cord contains**

- a) Vas deference**
- b) Artery to the vas & testicular artery**
- c) Pampiniform plexus of veins & accompanying lymphatics**
- d) Sympathetic plexus of nerves along the arteries.**

## **TESTICULAR PHYSIOLOGY**

**The two main function of the testis are to produce spermatozoa and to secrete testosterone. The seminiferous tubules and sertoli cells are responsible for spermatogenesis and the interstitial cells or Leydig cells secrete androgen, predominantly testosterone.**

**Specialised functional complex between sertoli cells are believed to form the bloodtestis barrier. FSH is required to initiate spermatogenesis. Between the seminiferous tubules of the testis lies the interstitial tissue containing blood vessels, lymphatic vessels, Leydig cells, macrophages and supporting cells. The leydig cell produces androgen, mostly testosterone, under the influence of LH.**

# **DIAGNOSTIC APPROACH TO SCROTAL SWELLINGS**

## **CLINICAL EXAMINATION**

Some clinical findings are important to particular cases such as follows.

|                   |   |                                |
|-------------------|---|--------------------------------|
| Hydrocele         | - | Transillumination Positive     |
|                   | - | Able to get above the swelling |
|                   | - | Fluctuation Positive.          |
| Testicular tumour | - | Loss of testicular sensation   |
| Torsion Testis    | - | Prehn's sign Positive          |

## **GENERAL INVESTIGATIONS**

Routine haematologic investigations like Total WBC count, Differential Leucocyte count & ESR will be helpful in diagnosing inflammatory conditions causing scrotal swellings. Eosinophil count will be raised in Filariasis which is a common cause for secondary hydrocele in endemic areas. Night smear examination will reveal microfilariae in cases with filarial aetiology.

Urine examination and urethral discharge analysis will be helpful in probing into the causes of epididymo-orchitis. Gram staining and culture will be helpful not only in identifying the causative factor but also in testing specific antibiotic sensitivity.

Serologic studies will be of help in diagnosing specific cause like syphilis, HIV and nonspecific causes like Chlamydial infection.

Accessory investigation like slit smear exam for leprosy, hypersensitive test for tuberculosis & leprosy and pus culture from purulent lesions of the scrotal wall also help in moving towards an appropriate diagnosis.

### ***RADIOLOGICAL INVESTIGATION:-***

#### ***X- RAY CHEST***

Plain x- rays of chest may be helpful in further characterising the pathology by the evidence of pulmonary tuberculosis, eosinophilia or secondaries.

#### ***ULTRASOUND***

**USG Examination using high resolution transducers (5 to 7.5Mhz)** has now become the investigation of choice in scrotal swellings. Major diagnostic role of ultrasound is to differentiate intra and extra testicular lesions.

The aphorism that focal poorly reflective lesions within the testis is presumed malignant until proved otherwise, hold true. It help us in differentiating simple hydrocele from closely clinically mimicking conditions like cyst of epididymis and testicular tumor. Extreme value of ultrasound lies in diagnosing rupture of testis in case of trauma.

Entire genitourinary system, Liver and retroperitoneal lymph nodes could be simultaneously evaluated.

### ***DOPPLER***

It helps us in differentiating testicular torsion from acute epididymorrchitis by the flow void sequences in the testicular artery (Levy & Pederson et al 1998). The disadvantage of Doppler lies in non visualization of the testicular infarct produced in cases where spontaneous detorsion has occurred now.

### ***LYMPHOGRAPHY***

Bipedal Lymphography is now rarely performed as a staging procedure while it can detect metastasis in normal sized node and the results of lymphography are equivocal to abdominal CT. this invasive procedure is not reliable in detecting recurrence compared to the use of CT and tumor markers.

### ***ODILAG (OPEN DIRECT INGUINAL LYMPHNGIOGRAPHY)***

It was preferred over by pedal lymphangiography due to high percentage of false negative results because of poor filling of primary lymph nodes in the later case. On the table ODILAG is helpful in identifying overlooked diseased nodes. This modality is now not in vogue due to the complications like Ethiodol pneumonitis and the advent of CT.

### ***COMPUTED TOMOGRAPHY***

CT is the mainstay in the radiological staging of testicular tumors. CT. helps in the identification of nodes between 8 to 10mm size. Other abdominal organs can also be scanned at the same time. An initial staging thoracic CT is indicated as it is the most sensitive methods in detecting pulmonary and

mediastinal metastasis compared to lymphography. CT is a non invasive procedure and also readily identifies the sentinel node and thence picking out early metastasis. The disadvantage being misdiagnosis of unopacified bowel loops, lymphatic vessels, post operative haematomas as lymphadenopathy and thereby overstaging. An added advantage of CT being its guidance in obtaining specimens from retroperitoneal lymph nodes.

### ***MRI***

MRI has its role in diagnosing residual tumor from retroperitoneal fibrosis which is very difficult to diagnose by CT alone. The disadvantage of MRI is failure in detecting calcification. The major disadvantage being unavailability, time consumption and cost factor.

### ***RADIONUCLEOTIDE SCANNING***

<sup>99m</sup>Tc Technetium pertechnetate scan is a widely practiced, accepted procedure for diagnosing testicular torsion (Heck et al 1974). The disadvantage being cost effectiveness and availability.



## ***RESULTS OF STUDY AND DISCUSSION***

During the two years of study 324 cases were admitted in the Department of General surgery, GRH, MMC with swellings of scrotum, of these 239 were primary hydroceles of various types making about 74% of the total. The other common causes we came across in this study were epididymoorchitis and varicocele both accounting for about 14%. The remaining 12% were due to testicular tumor, Fourniers gangrene, epididymal cyst, scrotal abscess, haematocoeles, spermatoceles and sebaceous cyst as shown below.

Total number of scrotal swellings admitted during the period of June – 04 to September –06.

|    |                    |     |        |
|----|--------------------|-----|--------|
| 1  | Primary Hydrocele  | 239 | 73.76% |
| 2  | Epididymoorchitis  | 27  | 8.33%  |
| 3  | Varicocele         | 18  | 5.55%  |
| 4  | Sebaceous Cyst     | 12  | 3.71%  |
| 5  | Testicular Tumour  | 11  | 3.39%  |
| 6  | Fourniers Gangrene | 6   | 1.85%  |
| 7  | Epididymal Cyst    | 4   | 1.23%  |
| 8  | Heamatocele        | 3   | 0.92%  |
| 9  | Scrotal Abscess    | 2   | 0.61%  |
| 10 | spermatocele       | 1   | 0.30%  |
| 11 | Torsion testis     | 1   | 0.30%  |

## ***HYDROCELE***

Hydrocele is an abnormal collection of serous fluid in the vaginalis of the testis or within some part of processus vaginalis.

It can be Congenital or Acquired and Primary (idiopathic) or secondary.

It is better defined as abnormal collection of serous fluid in tunica vaginalis when no other pathology is made out in the testis or epididymis.

Hydrocele is the commonest of all scrotal swelling that we encountered in our series we came across 239 cases of primary hydrocele and 9 cases of secondary hydrocele.

|                  | <b>No</b> | <b>Percentage</b> |
|------------------|-----------|-------------------|
| <b>Primary</b>   | 239       | 96.37%            |
| <b>Secondary</b> | 9         | 3.36%             |
| <b>Total</b>     | 248       | 100%              |

### ***I. Primary Hydrocele***

Types of primary hydrocele are

- a. Vaginal (Commonest)
- b. Infantile
- c. Congenital
- d. Funicular
- e. Encysted hydrocele of the cord

***Other rare types are***

- a. Hydrocele of the hernial sac
- b. Hydrocele en bisac
- c. Hydrocele of canal of Nuck

## ***AETIOPATHOLOGY AND MANAGEMENT***

### ***AETIOLOGY***

The various aetiologies proposed are excessive production of fluid, defective absorption, interference with lymphatic drainage and connection with a hernia of the peritoneal cavity.

### ***CLINICAL FEATURES***

Usually fluctuant, not reducible and it will be possible to get above the swelling.,

#### ***a. VAGINAL HYDROCELE***

This is the commonest variety commonly found in the tropical countries and the processus is found obliterated at the level of superior pole of the testis. The fluid is amber coloured and the specific gravity ranges from 1.022 to 1.024. The usual composition is water, inorganic salts and 6% albumin; fibrinogen is present in longstanding cases and cholesterol and tyrosine crystals can be found rarely.

It starts as a painless swelling in a middle aged, man, often unilateral and the testis cannot be felt separately.

***Pattern of age distribution in primary vaginal hydrocele***

Based on our study the pattern of age distribution in patients with primary vaginal hydrocele is given below. The maximum number of patients presented during 31-40 years of age.

| S. No.       | Age group | No. of pts | Percentage  |
|--------------|-----------|------------|-------------|
| 1            | 11-20yrs  | 17         | 7.4%        |
| 2            | 21-30 yrs | 39         | 17.2%       |
| 3            | 31-40 yrs | 74         | 32.6%       |
| 4            | 41-50 yrs | 49         | 21.6%       |
| 5            | 51-60 yrs | 30         | 13.2%       |
| 6            | >60 yrs   | 18         | 7.9%        |
| <b>Total</b> |           | <b>227</b> | <b>100%</b> |

**b. CONGENITAL HYDROCELE**

**syn: COMMUNICATING HYDROCELE**

It is due to patent processus vaginalis, having direct communication with the peritoneal cavity; but the orifice at the deep inguinal ring is too small to develop hernia. It presents since birth and slowly disappears on lying down position.

Ascities or Ascitic tuberculous peritonitis should be considered if the swellings are bilateral.

### **c. INFANTILE HYDROCELE**

Here the tunica and the processus vaginalis are distended upto the deep inguinal ring but do not communicate with the general peritoneal cavity. It presents as a inguinoscrotal swelling not reducible and there will be no impulse on coughing. It does not disappear on lying down.

### **d. FUNICULAR HYDROCELE**

Rare condition often confused with inguinal hernia. The funicular process is closed just above the tunica vaginalis so that it does not produce a proper scrotal, but an inguinal swelling is produced. Testis can be felt separately.

### **e. ENCYSTED HYDROCELE OF THE CORD**

It occurs when a portion of the funicular process persists and remains patent, shut off from the tunica vaginalis below and the peritoneal cavity above. It starts as an oval cystic swelling in relation to the spermatic cord lying in the inguinal / inguinoscrotal / scrotal region. Testis can be felt separately not reducible and it will be possible to get above the swelling. Traction test is pathognomonic.

### **f. BILOCULAR HYDROCELE**

**syn: HYDROCELE EN BISAC**

Rare variety where the hydrocele has intercommunicating sacs, one above and one below the neck of the scrotum. The upper sac has no

connection with processus vaginalis and it is infarct the herniated tunica vaginalis. Cross fluctuation between the two sacs of the hydrocele is pathognomonic.

#### **g. HYDROCELE OF THE HERNIAL SAC**

Due to stagnation of the fluid within the hernial sac when a tag of omentum blocks the opening of the sac at the deep inguinal ring.

Among the primary hydrocele cases primary vaginal hydrocele was the commonest type in our series. The other types like congenital, infantile and encysted hydrocele of the cord were found rare. No funicular hydrocele case was come across during the period of study.

| <b>Type</b> | <b>No.</b> | <b>Percentage</b> |
|-------------|------------|-------------------|
| Vaginal     | 227        | 94.9%             |
| Encysted    | 6          | 2.5%              |
| Congenital  | 3          | 1.2%              |
| Infantile   | 3          | 1.2%              |
| Total       | 239        | 100%              |

## **II. SECONDARY HYDROCELE**

It is due to the disease of the testis and or the epididymis.

Causes of secondary hydrocele are acute conditions like acute epididymoorchitis, torsion, trauma and chronic conditions like chronic epididymoorchitis, malignant disease of the testis, lymphatic obstruction and syphilitic affection of the testis.

These are usually small and lax and hence testis is easily palpable. Any acutely developing hydrocele may be secondary to testicular tumor.

In our study the major cause of secondary hydrocele is by epididymoorchitis, followed by testicular tumor.

| <b>Cause</b>         | <b>No.</b> | <b>Percentage</b> |
|----------------------|------------|-------------------|
| 1. epididymoorchitis | 7          | 77.8%             |
| 2. Testicular Tumor  | 2          | 22.2%             |

## **COMPLICATIONS OF HYDROCELE**

### **a. INFECTION**

### **b. ATROPHY OF THE TESTIS**

Dey has observed an arrest of spermatogenesis and consequent testicular atrophy as a result of fluid pressure in the tunica vaginalis.

### **c. IMPAIRED SPERMATOGENESIS**

M.C. Dandapat et al have reported that big hydroceles of the tunica vaginalis testis of long duration impair spermatogenesis or fertility.

M.C.I.

- d. **RUPTURE** – traumatic or spontaneous
- e. **HAEMATOCELE** – traumatic or spontaneous
- f. **HERNIA OF THE HYDROCELE SAC**
- g. **CALCIFICATION OF THE HYDROCELE SAC WALL**
- h. **PYOCELE**
- i. **CLOTTED HYDROCELE**

Most of the primary hydroceles were simple except for a few which were complicated.

|             |     |        |
|-------------|-----|--------|
| Simple      | 226 | 94.56% |
| Complicated | 13  | 5.44%  |
| Total       | 239 | 100%   |

The commonest complication we came across were pyocele and haematocele. Other complications like calcification of the sac and herniation of the hydrocele sac were rarely met.

| <b>Complications</b>           | <b>No.</b> | <b>Percentage</b> |
|--------------------------------|------------|-------------------|
| 1. Pyocele                     | 6          | 46.2%             |
| 2. Haematocele                 | 5          | 38.5%             |
| 3. Calcified sac               | 1          | 7.7%              |
| 4. Herniation of hydrocele sac | 1          | 7.7%              |
| Total                          | 239        | 100%              |



All the hydrocele patients included in this study had a total and differential count done as a routine investigation. 37 patients were found to have eosinophil count more than 5%.

## **TREATMENT MODALITIES AVAILABLE FOR HYDROCELE**

### **a. ASPIRATION**

It is preferred in older patients who are unfit for surgery. Before aspiration position of the testis should be confirmed and ensured that it is healthy.

Disadvantages of this procedure are it gives only temporary relief and rarely it causes haematoma.

### **b. SCLEROTHERAPY**

The solution which was used in the earlier days is Quinine hydrochloride (4gm) and Urethane (4gm) in water (30ml).

Sclerosants more commonly used now are Sodium tetradecylsulphate, Tetracycline hydrochloride, Minocycline, Ethanolamine oleate and Polidocanol (3%).

After aspirating the fluid sclerosant is injected and the scrotum is supported for some days. If necessary the procedure may be repeated at a later date.

Disadvantages are pain, recurrence, haematoma and infection.

Various studies have been conducted on sclerosants for hydrocele using various sclerosants which includes phenol (Nash 1984), tetradecyl sulphate (Mecfarlane 1983). Tetracycline (Hu et al, Badenech et al, Bullock et al), Ethanolamine oleate (Hell storm et al) and Polydocand (Fuse et al). Upon the above sclerosants comparative trial by Rencken et al revealed that Tetracycline + Tetradecyl sulphate to be superior in the cure rates (95%), availability and also cost effectiveness.

### **c. SURGICAL MANAGEMENT**

There are two approaches – inguinal and scrotal. Both procedures can be done under LA, SA or GA.

In inguinal approach the sac is delivered through an inguinal incision over superficial inguinal ring.

In the scrotal approach the sac is delivered through vertical paramedian or transverse incision.

If the hydrocele is small and thin walled Jaboulay's Procedure or Lord's Plication can be done. If the hydrocele is large or the sac is thick walled, excision of the sac is the procedure of choice. If hematoma formation is expected scrotum should be drained by a CRD. Risk of post operative complications is less after Lord Procedure and it can be done as an out patient procedure. (Singh. Dr et al)

In our 2 years of study 248 surgeries were performed on patients presented with hydrocele. The selection of the patient for the type of surgery was based upon size, thickness of the tunica vaginalis, redundancy of the scrotal skin and also the surgeon who is performing the procedure. Smaller sacs were either everted and plicated and the larger ones were excised. In some cases both partial excision and eversion were performed. Very few patients with increased redundancy of scrotal skin underwent scrotal skin excision and reconstruction following usually excision of sac.

The statistical data of the procedure adopted in our patients are as follows. Surgeries done for bilateral hydrocele are considered as two separate surgeries.

| <b>Procedure</b>               | <b>No.</b> | <b>Percentage</b> |
|--------------------------------|------------|-------------------|
| 1. Eversion                    | 112        | 45.1%             |
| 2. Excision                    | 82         | 33.0%             |
| 3. Partial Excision & Eversion | 15         | 5.9%              |
| 4. Lord's plication            | 27         | 11.8%             |
| 5. Orchiectomy                 | 9          | 3.6%              |
| 6. Herniotomy                  | 3          | 1.2%              |
| <b>Total</b>                   | <b>248</b> | <b>100%</b>       |

## INCISIONS

Vertical para median rapheal incision was the more common incision used in our subjects. Transverse incision was used for small and medium sized hydroceles accounting for about 13% of cases.

| <b>Incision</b> | <b>Percentage</b> |
|-----------------|-------------------|
| 1. Vertical     | 92%               |
| 2. Transverse   | 8%                |
| <b>Total</b>    | <b>100%</b>       |

## POST OPERATIVE COMPLICATIONS

### 1. INFECTION

### 2. SCROTAL HEAMATOMA

Most of the patients had an uneventful post operative period. The commonest complications we encountered were post operative infection and post operative haematoma.

|               |    |        |
|---------------|----|--------|
| 1. Infections | 27 | 10.88% |
| 2. Heamatoma  | 9  | 3.62%  |

## COMPARISON OF COMPLICATIONS OF VARIOUS SURGICAL PROCEDURES

| SURGICAL PROCEDURE          | Total No. | INFECTIONS |       | HAEMATOMA |      |
|-----------------------------|-----------|------------|-------|-----------|------|
|                             |           | No         | %     | No.       | %    |
| Excision                    | 82        | 13         | 15.8% | 6         | 7.3% |
| Eversion                    | 112       | 11         | 9.8%  | 2         | 1.8% |
| Partial Excision & Eversion | 15        | 2          | 13.3% | 1         | 6.6% |
| Lord's Plication            | 27        | 1          | 3.7%  | 0         | 0%   |

## OTHER NEW PROCEDURES UNDER TRIAL ARE

### a. EVERTED PLICATION (Rohondia OP et al)

It is a combination of plication and eversion. The advantages claimed are need of less time less suture material and less injury to epididymis and testis.

### b. ENDOSCOPIC HYDROCELE FULGRATION (Nishiyama et al)

This procedure is under evaluation. Here the partial surface of the sac is completely fulgurated by a resectoscope with a 30 degree lens inserted via a modified laparoscopic trocar inserted into the hydrocele under video monitoring.

### c. LAPAROSCOPIC REPAIR OF PAEDIATRIC HYDROCELES (Janetscheck. G et al)

Here the open processus vaginalis is transected using laparoscope.

## **VARICOCELE**

Varicocele is defined as dilated elongated and tortuous pampiniform plexus of veins.

### **Aetiology**

a) Primary or idiopathic

b) Secondary

Left – due to hypernephroma

Right – Tumor thrombus extending upto IVC resulting in block

c) Constitutional

It is more common in left side (98%) when compared to right side (2%). It is usually asymptomatic when small, but causes a heavy, dragging, aching sensation which becomes worse on prolonged standing and exertion. Some present with subfertility and infertility. Any middle aged men with sudden onset of varicocele must be completely investigated for hypernephroma.

On inspection dilated veins can be seen over the scrotum; on palpation the dilated veins will appear like a bag of worms. Cough impulse and thrill will be usually positive. Testis on the affected side is usually small due to atrophy. On lying down primary varicocele empties, whereas it does not happen in case of secondary lesion.

### ***PREDILICTION FOR LEFT SIDE***

1. High insertion and increase in length of the left testicular vein.
2. The left testicular vein drains into the left renal vein at right angle.
3. The left suprarenal vein drains into the left renal vein results in release of catecholamines causing vasoconstriction.
4. Compression of left renal vein in between abdominal aorta and superior mesenteric artery.
5. Compression by loaded rectum.
6. Left testicular artery may arch over left renal vein and thus may cause compression over it.

Varicocele accounted for 5.5% of all cases in our series. Majority of them were left sided (80%) and among the 18 cases infertility was reported in only one case.

| Left    | Right   |
|---------|---------|
| 14      | 4       |
| (77.8%) | (22.2%) |

All these cases were subjected to ultrasound examination. Multiple dilated tubular structure suggestive of varicocele were identified in scrotal ultrasound. None of the patients had any associated abdominal pathology.

## **TREATMENT OF VARICOCELE CONSERVATIVE**

If the varicocele is small scrotal support and reassurance is enough.

## **SURGICAL**

The indication for surgery are severe pain, infertility and for cosmetic purpose. For all cases seminal analysis was done pre operatively. Only one case was found to be infertile.

## **TYPES OF SURGERY CLASSICAL**

Here a portion of spermatic veins is excised between ligatures. It can be done through inguinal or scrotal route, scrotal approach is not usually preferred as recurrence, bleeding, infection and damage to testicular artery are more common.

## ***PALOMO OPERATION***

Here the spermatic vein is approached retroperitoneally and ligated.

All these 11 cases were operated upon. Ligation and excision was the procedure adopted for all these patients.



### ***LAPAROSCOPIC SURGERY***

Here the internal spermatic vein is clipped and divided. Laparoscopic varicocelectomy is a simple, safe, effective and minimally invasive procedure and it can be proposed as a viable alternative to open traditional surgical methods. Recurrence rate is very low when compared with traditional surgical methods (Recurrence rate for traditional methods is 1-25%) (Berolli et al). Patient can return to normal activity earlier.

### **MICROSURGICAL HIGH INGUINAL VARICICOCELECTOMY WITH DELIVERY OF THE TESTIS**

In this method the testis is delivered through inguinal route and the spermatic cord is dissected under optical magnification. All external spermatic and gubernacular veins are ligated and testicular artery, lymphatics and vas deferens is identified and preserved.

Advantages of this procedure are no clinical recurrence, no post operative hydrocele formation, increase pregnancy rate and no wound infection (Chaloothy E et al).

### **EMBOLISATION**

Embolisation of testicular vein at the inguinal canal is performed using either balloons or steel coils.

### ***PERCUTANEOUS SCLEROTHERAPY***

It is a simpler technique obliterates the testicular vein by placing the catheter tip at the proximal portion of the testicular vein and injection of sclerosants.

### ***EPIDIDYMOORCHITIS***

It refers to inflammation of the epididymis and the testis.

Clinically it is divided into acute and chronic epididymoorchitis; pathologically it is divided into specific and nonspecific epididymoorchitis.

Epididymoorchitis accounted for 8% of the cases in our series. 7 out of 27 patients presented with secondary hydrocele.

| Type                | No. | Percentage |
|---------------------|-----|------------|
| 1. Specific<br>(TB) | 8   | 29.6%      |
| 2. Nonspecific      | 19  | 70.4%      |
| <b>Total</b>        | 27  | 100%       |

### ***ACUTE EPIDIDYMOORCHITIS***

Initially the infection is confined to the epididymis and later spreads to the body of the testis. Infection can reach the epididymis in the following ways.

As retrograde infection from the urethra, prostate and seminal vesicles; globus minor is affected first. Organisms involved in retrograde infection are E.coli, Klebsiella and Gonococcus.

As sexual transmission associated with urethritis; organisms involved are Chlamydia and Gonococcus.

As blood borne infection; globus major is affected first. Organisms responsible are Streptococci, Staphylococci, Proteus and E.coli.

These patients present with severe pain and swelling in the testis and associated constitutional symptoms like fever, malaise and chills. Urinary tract infection symptoms like frequency, urgency, dysuria, pyuria and haematuria may be present. On examination the scrotal wall will be red, edematous, shiny and adherent to the epididymis. Epididymis will be diffusely swollen and tender.

### ***DIFFERENTIAL DIAGNOSIS FOR ACUTE EPIDIDYMOORCHITIS***

- a. Acute torsion of spermatic cord
- b. Acute torsion of hydatid of morgagni and other appendages of testis.
- c. Spontaneous haemorrhage of a testicular tumor
- d. Trauma to the scrotum
- e. Thrombosis of the pampiniform plexus

In our study 11 of the 27 patients with epididymo-orchitis made an acute presentation. All the patients were managed conservatively. Follow up proved the patients to be symptom free with the return of testis to the normal size showing no residual changes.

| <b>Signs &amp; symptoms</b> | <b>No. of pts</b> |
|-----------------------------|-------------------|
| Pain                        | 11                |
| Fever                       | 5                 |
| Swelling                    | 9                 |
| Urinary symptoms            | 2                 |

### ***TREATMENT OF ACUTE EPIDIDYMO-ORCHITIS***

In children and men over 35 years the infection is mostly due to bacterial urinary tract infection. So the urine should be cultured and appropriate antibiotics should be given for a period of 2-3 weeks.

In young men below 35 years the infection is mostly sexually acquired. So Gram stain and special culture of urethral secretion should be done and treated with appropriate antibiotics.

If culture is not helpful, serological tests like (IgM and IgG antibodies to Chlamydia) is done and treated with Tetracycline and Doxycycline. Minocycline and Erythromycin can also be used.

Non specific measures are absolute bed rest, plenty of oral fluids, elevation of scrotum and analgesics.

### ***ACUTE EPIDIDYMOORCHITIS OF MUMPS***

It occurs in prepubertal boys. It occurs in 20% of males suffering from Mumps as the parotid swelling is waning. It starts as unilateral testicular swelling with severe pain, high fever, malaise and associated with acute hydrocele. There will be no scrotal edema. If there is bilateral affection it can cause infertility. Partial atrophy will cause persistent testicular pain. As the cure is spontaneous symptomatic treatment alone is needed.

Other rare causes of acute epididymoorchitis are Enteroviral infection, Meningococcal infection, Brucellosis, Typhoid, LGV, Trauma, Exposure to chemicals and Blastomycosis.

### ***CHRONIC EPIDIDYMOORCHITIS***

It can be classified into specific and nonspecific epididymoorchitis. Specific infections are of Tuberculous, Syphilitic, Leprous and Viral etiology.

### ***CHRONIC TUBERCULOUS EPIDIDYMOORCHITIS***

Globus of the epididymis is the first part to be affected by retrograde infection from a Tuberculous focus which is usually in the seminal vesicle. Onset of this condition is insidious and it is rarely through a haematogenous route either from a focus in the lungs or from isolated renal tuberculosis.

Patients present with a minimal swelling of the testis and slight aching sensation. In 30% of cases there is an associated secondary hydrocele.

It starts as a firm discrete swelling in the lower pole which becomes firm and craggy in consistency. In the later stages nodules could be appreciated in the body which further softens to become cold abscess and discharging sinus in the posterior aspect of the scrotum. Characteristic beading of the vas deferens due to sub epithelial pearls, thickened seminal vesicles, tender and irregular prostate are the late manifestations of this disease.

## **TREATMENT**

If no significant improvement with antituberculous treatment is made out, epididymo-orchidectomy is usually performed.

In our series 8 cases have been reported to be of tuberculous etiology among the 27 epididymo-orchitis. 6 cases were clinically diagnosed and trial ATT was given which showed no clinical improvement which further made the patient to undergo low orchidectomy. The remaining 2 patients were misdiagnosed both clinically and sonologically as testicular tumor. Hence high orchidectomy was performed and the biopsy revealed the truth.

### ***SYPHILITIC ORCHITIS***

Testis and epididymis are affected both in congenital and acquired syphilis. Invariably the testis being affected first. Chronic draining fistulae are common in untreated conditions.

Manifestation of syphilis are of 3 types.

- a. **Bilateral orchitis** – in congenital syphilis
- b. **Interstitial fibrosis** – symptomless condition in which the testis of normal size but for the loss of testicular sensation, and is usually bilateral.
- c. **Gumma** – This is the commonest type which is always unilateral. It starts as a painless, hard, enlarged testis with loss of testicular sensation. Later the testis softens anteriorly and a gummatous ulcer develops in the anterior aspect of the scrotal wall. Secondary hydrocele is a must in this condition and is to be differentiated from testicular tumor.

### ***LEPROUS ORCHITIS***

In case of lepromatous orchitis the testis is usually extensively involved with destruction of seminiferous tubules eventually resulting in testicular atrophy and consequent sterility.

No case of leprous orchitis was reported in this series.

### ***VIRAL ORCHITIS***

Usually presents as an acute manifestation and may sometimes persist chronically with reactive secondary hydrocele. The common causative agents being Mumps virus, Influenza, Smallpox, Measles, Varicella, Coxsackie virus and HIV.

### **TREATMENT**

Symptomatic treatment with analgesics and steroids. ACTH is rarely used and is less useful. Sometimes tapping of a tense hydrocele is done to relieve pain but with risk of secondary infection.

### **CHRONIC NONSPECIFIC EPIDIDYMOORCHITIS**

#### **(GRANULOMATOUS / AUTOIMMUNE)**

It occurs in middle aged men following an acute attack which fails to resolve. It often presents with fever and a moderately tender testicular swelling. This condition is to be differentiated from tuberculous epididymoorchitis by its larger size and smoothness.

It usually, follows Sarcoidosis, chemical exposure, urethral stricture causing reflux of urine down the vas initiating an autoimmune reaction.

### **TREATMENT**

Epididymo-orchidectomy.



### ***POST OPERATIVE EPIDIDYMOORCHITIS***

In earlier days post operative epididymoorchitis following trans vesical prostatectomy was frequently encountered. Vasectomy was employed for the above cases to prevent spread of infection which now seldom occurs, after the invention of modern chemotherapeutic agents.

### **SUB ACUTE EPIDIDYMOORCHITIS (FILARIAL)**

If filarial epididymoorchitis the globus major is the first part to be affected followed by testis. In the initial stages the digital fossa between the testis and the epididymis on the lateral aspect is obliterated. Later the testis enlarges and becomes firm with loss of testicular sensation.

No case of the above condition was reported in our series.

Urine culture was done for the clinically suspected 27 patients with epididymoorchitis, 9 of the 27 turned out to be positive. The details of the results are given below.

| <b>Organisms grown</b> | <b>No.</b> |
|------------------------|------------|
| 1. E.Coli              | 6          |
| 2. Klebsiella          | 3          |

## ***TREATMENT***

### ***A. MEDICAL***

- DEC 100 mgm 3 times a day × 2-3 weeks
- Paramelaminy1 Phenyl Stilbonate which acts on the infective larvae and the immature adult worms.
- Antibiotics (Aminoglycosides)

### ***B. SURGICAL***

#### **Orchidectomy (low)**

The patients who were diagnosed clinically as epididymoorchitis were treated conservatively except for 8 cases of tuberculous epididymoorchitis, and an other testicular abscess who underwent low orchidectomy.

1. Conservative      25

2. Orchidectomy      2

## ***TORSION OF THE TESTIS***

It is defined as torsion of the cord with characteristic rotation of the testis and an anterior presentation of the testis. Torsion is relatively uncommon in fully descended testis as it is well anchored and can not rotate.

## **AETIOLOGY**

- a. Inversion of testis
- b. Long mesorchium
- c. High inversion of tunica vaginalis
- d. Undescended or ectopic tests

- e. Voluminous tunica vaginalis
- f. Separation of tunica vaginalis from testis

## **TYPES**

### **A. INTRAVAGINAL (PRE PUBERTAL MALES)**

Occurs due to long mesoorchium and torsion occurs within the tunica vaginalis.

### **B. EXTRAVAGINAL TORSION (NEONATES)**

When the cord twists above the mesoorchium the whole tunica rotates resulting in vascular occlusion and further gangrene of testis within 24 hours. It is unrelated to anomalous suspension of testis.

Intravaginal torsion has a higher incidence when compared with extravaginal torsion (Skogland et al 1970).

Torsion usually follows straining, lifting heavy objects or during coitus due to violent contractions of cremaster. Both the testis rotates in a fashion so as to face the midline. Torsion presents as severe pain in the testis or groin referred to the lower abdomen associated with nausea or vomiting. The scrotum is swollen and edematous within 2 to 3 hours and the testis and epididymis lie is swollen and edematous within 2 to 3 hours and the testis and epididymis lie in an abnormal position in the high or horizontal in addition of positivity of Prehn's sign. Surgical restoration of blood flow within 5 hours salvages the testicular function in 83% of patients (Skogland et al).

### ***TREATMENT***

- a. Within 3-4 hours of the episode manual derotation should be tried by gentle manipulation.
- b. Exploration of testicular torsion – untwisting of the testis, trimming of the excessive tunica vaginalis with fixation of the testis to the lateral wall or high orchidopexy is performed. The same procedure is performed to the opposite side as the predisposing congenital abnormality to this conditions bilateral.
- c. If the patient presents after complete jeopardisation of the testicular tissue diagnosed pre or per operatively removal of the infarcted testis is done.

### ***TORSION OF TESTICULAR APPENDAGES***

The torsion of testicular appendage results in a palpable localized painful lump on the upper pole of the testis (Skogland et al 1970). Transillumination reveals a characteristic “BLUE DOT SIGN” which is the infarcted appendage (Holland et al 1981).

### ***TREATMENT***

If the diagnosis is certain conservative management is all sufficient (Holland et al 1987). If any doubt exist exploration followed by excision is advised. In our two years of study one testicular torsion case was reported and which was treated by orchidectomy and contralateral orchidopexy.

## **HAEMATOCELE**

It is defined as haemorrhagic effusion into the tunica vaginalis, it is broadly classified into recent onset and old haematocele.

### ***A. RECENT HAEMATOCELE***

Commonly presents with a history of trauma followed by severe pain, tenderness and absence of transillumination. Other causes being surgical procedures like tapping of hydrocele or following vasectomy. Rarely tumors are said.

### **TREATMENT**

Exploration of tunica with evacuation of clots. If testis is ruptured in a linear fashion careful suturing is advised; for a segmental damage wedge resection is usually employed.

### ***B. OLD HAEMATOCELE***

It is due to slow bleed into the tunica and hence painless. Closely mimics a tumor but can be differentiated by means of testicular sensation.

### **TREATMENT**

It is again surgical exploration and if testis is atrophied or low orchidectomy is employed.

## **RESULTS**

Of the 324 subjects included in our study 8 cases were proved to be haematocele. 6 cases of haematocele were due to complication of primary hydrocele.

Among the 8 patients 5 were subjected to low orchidectomy as the testis were atrophied and the remaining underwent testis preserving procedures.

## **SPERMATOCELE**

It is defined as a retention cyst which arises from vasa efferentia or from the sperm conducting system of the epididymis. It contains fluid of barley water appearance rich in spermatozoa. It presents as a small, painless, soft fluctuant swelling situated at the head of epididymis.

## **TREATMENT**

Smaller ones are self resolving, the larger needs either aspiration or excision. Sclerotherapy with Quindine hydrochloride / Urethane or with Tetradecyle can also be tried.

Only one case was reported in all these two years and was diagnosed pre operatively by ultrasound which showed an anechoic lesion containing turbid fluid. The cyst was excised and the contents proved the earlier diagnosis.

### ***SPERM GRANULOMA***

Escape of spermatozoa in the epididymis results in this painful swelling. It is commonly seen after vasectomy.

### ***TREATMENT***

Excision

No case was reported in the years of our study

### ***CYST OF EPPIDIDYMIS***

Multilocular crystal clear fluid filled cyst representing cystic degeneration of epididymis commonly presents in the middle aged men. They are thought to be remnants of mesonephric or paramesonephric ducts. The cluster of thin walled cysts resembling bunch of grapes are felt separately from the testis.

### ***TREATMENT***

Excision

In our series 4 cases were reported forming 1.2% of the total. All the 4 were treated by excision.

### ***SUBCUTANEOUS EMPHYSEMA –SCROTUM***

The above condition results commonly following chest injury. It is due to tracking of air in the subcutaneous tissue of the scrotum.

In our study no case was reported.

### ***SCROTAL ABSCESS***

It results due to infection of the scrotal skin as that of any other of the human body. Commonest causative agent is staphylococcus aureus. Diabetics are prone for the above condition.

In our series 2 cases were reported and 1 of them happened to be diabetics. All the cases were treated by incision and drainage and the pus sent for culture showed positively for staphylococcus in both cases.

### ***CARCINOMA SCROTUM***

Squamous cell carcinoma of the scrotum usually an occupational hazard takes the form of

Wide excision with en bloc dissection of inguinal nodes.

In our series no case of carcinoma scrotum was reported.

### ***FOURNIERS GANGRENE***

Abrupt rapidly progressive gangrenous infection of the genitalia which was earlier thought to be idiopathic is now given a definitive etiology (Gray 1960, Flanigan et al 1978), (Lamb and Juler 1983), (Carol et al 1986) traced the origin of infection from rectum, GUT or the skin and thereby disproved the definition given by Fournier as idiopathic scrotal edema in the year 1883.



The commonest causative organism being *Clostridium welchii*, haemolytic streptococci, *E.Coli*, *Bacteroides* and *staphylococcus aureus*. The predisposition for patients with diabetes is well documented and the reason behind being diabetic vasculopathy, defective phagocytosis and functional urinary tract obstruction form diabetic neuropathy (Yhornton 1971).

3 of our patients were diabetics and non of our 6 cases had rectal foci of infection. Urine culture revealed positive results in 4 of our 6 patients.

| <b>E.coli</b> | <b>Staph. aureus</b> |
|---------------|----------------------|
| 3             | 1                    |

## **TREATMENT**

The treatment begins with surgical debridement immediately upon diagnosis, with exploration within 24 hours to confirm the arrest of infection. Broad spectrum antibiotics directed at both aerobic and anaerobic bacteria should initiated immediately awaiting culture results. In most cases out standard regime was to start the patient of Triple Antibiotic coverage with Ampicillin, Gentamycin and Metronidazole. Newer Cephalosporins are used based on the investigation results. Despite extensive infection and widespread debridement the redundancy of scrotal skin allows compete or partial reconstruction by delayed closure. Larger or residual defects are managed with skin grafts. The skeletonised testicles following debridement are protected in

thigh pouches on a interim or long term basis. Orchidectomy is restricted to older patients with extensive surrounding tissue damage.

## **RESULTS**

In our series 6 cases of Fourniers gangrene were reported. Among them 4 were managed by secondary suturing, the remaining patients ere benefited with split skin grafts. Despite aggressive treatment morbidity and mortality remain high especially in older patients with rectal foci.

# TESTICULAR TUMORS

**Testicular tumor gains its importance as the most common tumor in young adult men between ages of 19 and 35 years. Modern chemotherapeutic regimens, meticulous nerve sparing RPLND, advanced sensitive radioimmunoassay for tumour markers, miraculous development in the imaging field has made us to know about the pathogenesis and outcome of our therapeutic trails. Neoplasms of the testis are always almost malignant, the exception being rare fibromas of the tunica vaginalis and pure leydig cell tumours.**

## MALIGNANT TUMOURS

Cancer of the testis comprises morphologically and clinically diverse group of neoplasms. The over whelming majority are primary and most of these are germ cell tumours. The management of each neoplasm is dependent on the histology and is influenced by the lymphatic and vascular drainage of the testis.

## **HISTOLOGICAL CLASSIFICATION**

|                                 |      |
|---------------------------------|------|
| SEMINOMA                        | 40%  |
| TERATOMA                        | 32%  |
| COMBINED SEMINOMA AND TERATOOMA | 14%  |
| INTERSTITIAL TUMORS             | 1.5% |
| LYMPHOMA                        | 7%   |
| OTHER TUMOURS                   | 5.5% |

## **INCIDENCE**

About 99% of testicular neoplasms are malignant, and though they make up only about 1 to 2 percentage of malignant tumours in men they are of the commonest forms of cancer in the young males (GLIBERT and HAMILTON).

In our study no particular age group was found to be more prone for testicular malignancy.

The histopathologic types encountered and the age incidence are given below.

| TUMOURS      | 20-29 | 30-39 | 40-49 | 50-59 | 60-69 | 70-79 | Total | %     |
|--------------|-------|-------|-------|-------|-------|-------|-------|-------|
| SEMINOMA     | 1     | 4     | 1     | -     | 2     | 1     | 8     | 72.7% |
| TERATOMA     | -     | -     | 1     | -     | -     | -     | 1     | 9.1%  |
| EMERYONAL    | 1     | -     | -     | -     | -     | -     | 1     | 9.1%  |
| CELL Ca      |       |       |       |       |       |       |       |       |
| NON HODGKINS |       |       |       |       |       |       |       |       |
| /B-CELL      | -     | -     | -     | -     | -     | 1     | 1     | 9.1%  |
| LYMPHOMA     |       |       |       |       |       |       |       |       |

In our series 11 cases of testicular tumours were reported making about 3.4% of the total. All the tumors were unilateral and most of them were left sided. 9 cases presented as painless testicular swelling and the remaining 2 were associated with testicular pain. Loss of testicular sensation was seen in almost all cases and secondary hydrocele was present in about 2 persons.

## **SEMINOMA**

Seminoma accounts for approximately 50% of all GCT's and most frequently appears in the fourth decade of life. Seminomas are uniform in gross and histologic appearance and are characterized by slow growth and late invasion. Metastatic spread is through the testicular lymphatics and dominates in the iliac, aortic and renal hilar nodes.

The typical or classic form consists of sheets of large cells with abundant cytoplasm and round hyperchromatic nuclei with prominent nucleoli. A lymphocytic infiltrate or granulomatous reaction with giant cells or both is frequently present. In the atypical form lymphocytic infiltration and granulomatous reactions are absent and necrosis is more common. Spermatocytic Seminoma is a rare histologic variant seen almost exclusively in men above the age of 45. It does not express placental alkaline phosphatase. Metastatic potential is minimal.

## **NON SEMINOMATOUS GERM CELL TUMOURS**

Non Seminomatous histology comprises about 50% of all GCT s and most frequently presents in the third decade of life. Most tumours are mixed consisting of two or more cell types (mostofi).

### **EMBRYONAL CARCINOMA (MTA)**

**Embryonal carcinoma is the most undifferentiated cell type. It is usually thought to be the most common testicular tumour of childhood; invasion and metastases occur earlier in the course of the disease. Because of the relatively rapid growth haemorrhage and necrosis are common. Metastasis to the abdominal lymphatics and the lungs may occur as an early event.**

### **CHORIOCARCINOMA (MTT)**

It is one of the most malignant tumors known. It consists of both cytotrophoblasts and syncytiotrophoblasts. Pure Choriocarcinoma is an extremely rare presentation. It often produces human chorionic gonadotrophin. The tumor is extremely rapidly invasive; metastasis may be both blood borne and through lymphatics and has usually occurred by the time of diagnosis. The prognosis for these patients is usually for worse than for patients because of

the advanced stage at the time of diagnosis. Elements of choriocarcinomas are frequently found in mixed tumours.

## **TERATOMA**

Teratoma arises from Totipotent cells in the rete testis and often contains a variety of cell types of which one or more predominate.

### **MATURE TERATOMA (TD)**

It consists of adult type differentiated elements, but cannot be considered as benign because such growths have metastasis.

### **IMMATURE TERATOMA**

It generally refers to tumour with partial somatic differentiation, similar to that seen in a fetus.

## **TERATOMA WITH MALIGNANT TRANSFORMATION**

It refers to a form of teratoma in which one of its components either immature or mature, develops aggressive growth and historically resembles another malignancy. They usually take the form of sarcomas and less frequently carcinomas.



## **YOLK SAC TUMOUR**

This tumour mimics the yolk sac of the embryo and produces Alpha-fetoproteins. The cells may have a papillary, granular or solid appearance; and may be associated with schiller – duval bodies. Pure yolk sac historically is the most common histology found in childhood GCT.

## **SERUM TUMOUR MARKES**

### **ALPHA\_FETOPROTEIN :**

Its production is restricted to Embryonal cell carcinoma and endodermal sinus tumours. In patients with pure Seminoma elevated concentration of this reflects an undifferentiated nonseminomatous element.

### **HUMAN CHORIONIC GONADOTROPHIN HCG:**

Elevated serum concentration can be found in patients with pure Seminoma as well as those with NSGCT.

### **LACTATE DEHYDROGENASE:**

Increase in serum concentration are a reflection of tumour burden, growth rate, proliferation and death.

## ***INITIAL PRESENTATION & MANAGEMENT***

### **SYMPTOMS AND SIGNS**

The pathognomonic presentation of a testicular tumour is a painless testicular mass that may range in size from a few mm to several cms. Only a minority of patients presents with painless mass; majority present with more diffuse testicular pain, swelling, hardness or some combination of these findings. Delay in diagnosis caused by their patient or physician related factors or both generally results in higher stage of presentation and presumably lower survival.

In no other disease testicular sensation is lost so early or so completely. Ten percent have a lax secondary hydrocele. Between 1 and 5 percent have gynaecomastia.

Atypical cases may simulate epididymoorchitis.

### **THE HURRICANE TUMOUR**

It is a ferocious malignancy which kills in a matter of weeks.

11 cases of testicular tumours were studied in our series making about

3.4

of the total. The various clinical presentations were as follows:

| <b>Symptoms / Signs</b>                    | <b>Seminoma</b> | <b>Teratoma</b> | <b>Embryonal Cell Ca</b> | <b>Lymphoma</b> | <b>Total</b> | <b>%</b> |
|--|-----------------|-----------------|--------------------------|-----------------|--------------|----------|
| <b>Painless, Heavy Testicular Swelling</b> | 8               | -               | 1                        | -               | 9            | 81.8     |
| <b>Painful testicular Swelling</b>         | -               | 1               | -                        | 1               | 2            | 18.2     |
| <b>Secondary Hydrocele</b>                 | 1               | 1               | -                        | -               | 2            | 18.2     |
| <b>Para aortic Lymphadenopathy</b>         | -               | -               | 1                        | -               | 1            | 9.1      |

**Two cases of the above seven presented with secondary hydrocele.**

**The classical sign of loss of testicular sensation was seen in all the patients.**

## DIAGNOSIS

A radical inguinal orchidectomy using an inguinal incision with early high ligation of the spermatic cord at the deep inguinal ring, minimizes local tumour recurrence and aberrant lymphatic spread and is the only acceptable therapeutic and diagnostic procedure.

In the rare situation where the diagnosis is in question, then an inguinal incision is required for an open biopsy.

Most of the cases were diagnosed clinically and relevant investigations were done to rule out other causes and to confirm the diagnosis. USG of testis and abdomen was done for all cases and the findings were as follows.

| USG Findings                | Seminoma | Terato Ca. | Embryonal cell Ca. | Lymphoma |
|-----------------------------|----------|------------|--------------------|----------|
| Homogenous echoic mass      | 6        | -          | -                  | -        |
| Mixed echogenic mass        | 2        | 1          | 1                  | 1        |
| Para aortic lymphadenopathy | 2        | 1          | 1                  | 1        |

Routine x-ray chest was taken for all cases. Mediastinal metastases were identified for the lymphoma patient.

### *STAGING*

A comprehensive evaluation is necessary to define the extent of disease and to determine the appropriate treatment and should include pathologic examination, determination of serum concentration of AFP and hcG and radiological Studies.

Broadly Stage I disease is confined to the testis; Stage II disease is restricted to the retroperitoneum (subdiaphragmatic) and stage III disease

represents involvement of supradiaphragmatic or other nodal status or visceral disease. Stage IV represents distant metastasis.

#### ***STAGING OF TESTICULAR TUMOUR***

Most of the tumours analyzed in this series were of stage I. Lymphoma testis was of stage III and the rest were stage II malignancies.

| <b>Tumour</b>             | <b>Stage I</b> | <b>Stage II</b> | <b>Stage III</b> |
|---------------------------|----------------|-----------------|------------------|
| <b>Seminoma</b>           | 7              | 1               | -                |
| <b>Teratocarcinoma</b>    | -              | 1               | -                |
| <b>Embryonal Cell Ca.</b> | -              | 1               | -                |
| <b>Lymphoma</b>           | -              | -               | 1                |
| <b>Total</b>              | 7 (63.6%)      | 3 (27.3%)       | 1 (9.1%)         |

Seven of the Seminomas presented with stage I disease and the remaining one seminoma presented with stage II disease. Both Teratoocarcinoma and Embryonal cell Carcinoma presned with stage II Disease, Lymphoma testis was of stage III.

#### **MANAGEMENT**

As soon as the diagnosis is arrived high inguinal orchidectomy (chevassou maneuver) was done.

## **CLINICAL STAGE I DISEASE : SEMINOMA**

The treatment options available are radiotherapy and observation.

### **RADIOTHERAPY**

Radiation therapy remains the treatment of choice. Dose is 150 to 180 cGY/day for five sessions per week. Total dose is 2500 to 3000 cGY. The relapse rate is negligible.

### **OBSERVATION**

Since long term follow up in needed observation is not considered routine.

### **NSGCT**

**The options available are RPLND, observation and chemotherapy.**

**As there is a predictable change lymphatic metastasis, modified bilateral**

**RPLND is usually the Conventional method of approach.**

**STAGE II DISEASE : NSGCT (Low burden tumour)**

The standard approach to patient's with clinical stage II A and some II B tumour has been RPLND. Margin of resection should not be compromised in an attempt to maintain ejaculatory Function.

## **SEMINOMA**

**Radiotherapy is the treatment of choice for most of the patients in this stage. The radiation portal and fraction is the same except that a boost of approximately 500-700 rads is administered.**

### **MANAGEMENT OF STAGE II & III DISEASE (High tumour Burden)**

Early Clinical trials developed regimens such as PVB (**cisplatin, vinblastine + Blemycin**) and VAB – 6 (**cisplatin, Vinblastine, Bleomycin, Dactinomycin + Cyclophosphamide**), eliminated maintenance therapy and replaced Vinblastine with Etoposide. Adjunctive Surgery was shown to be essential for achieving a disease free state. Although most patients were cured, significant adverse events were observed. So good and poor risk allocation algorithms were developed.

## **GOOD PROGNOSIS GERM CELL TUMOURS**

Good risk patients are those with a high likelihood of cure. Randomized trials permit a systemic evaluation of the least toxic maximally efficacious therapy for good risk patients.

#### **POOR RISK GERM CELL TUMOURS**

As the optimal CT regimen is yet to be determined, the patients coming under the above group should be managed by clinical trials. IT is done either by substitution or by dose intensification of an already existing one like **Vepeside + Holoxan + Adriablastine.**

#### **ADJUVANT CHEMOTHERAPY (STAGE II)**

Adjuvant CT remains a strong consideration in patients when six nodes or more are involved, any node is larger than 2cms or there is extra nodal extension. Two cycles of cisplatin based CT are nearly always effective. Etoposide has replaced vinblastine in adjuvant regimens. A recent study suggests that Etoposide plus cisplatin alone is adequate.

### **MANAGEMENT OF RESIDUAL DISEASE**

#### **RETROPERITONEUM:**

**NSGCT:-** There is general agreement over the need to resect all sites of measurable residual disease. A bilateral RPLND is required for residual NSGCT in retroperitoneum.

**SEMINOMA:-** observation is the choice of management for residual masses smaller than 3 cms. If more they are managed either by observation or surgical resection and direct immediate therapy.



## **MANAGEMENT OF RELAPSE AFTER CHEMOTHERAPY**

### **CONVENTIONAL DOSE SALVAGE THERAPY**

**Ifosfamide is combined with cisplatin and Etoposide (VIP) or cisplatin and Vinblastine (Velp) in patients whose disease is resistant to two prior regimens.**

### **HIGH DOSE THERAPY**

The drugs used in this regimen are **carboplatin** and **Etoposide** with or without an **oxazophosphorine (Cyclophosphamide or Ifosfamide)**.

### **NEW AGENTS**

A number of single agent trials have been conducted against refractory GCT. **Ifosfamide**, **Paclitaxel** and oral **Etoposide** have demonstrated antitumorous activity. Because **Paclitaxel** is synergistic with Cisplatin and **Oxasophosphorine** in vitro, it is being studied in combination with Cisplatin plus Ifosfamide.

### **ROLE OF SURGERY**

Surgery has curative potential in a highly selective group of patients with increase tumour marker level, even after Salvages CT.

### **BILATERAL TESTICULAR TUMOURS**

The incidence of bilateral testicular carcinoma varies between 0.5 to 7%. In general incidence metachronous testicular tumour is 4 times greater than that of synchronous variety (Christopher L. Coogen et al 1998) As there is 500-1000 times increase in risk of acquiring malignancy in the contralateral

testis in a patients with testicular tumour has resulted in performing biopsy of the contralateral testis and prophylactic radiotherapy for the same. Radiotherapy virtually eliminates the possibility of contralateral tumour, despite preserving its hormonal production but eliminates the fertility potential. Treatment is based on pathology and clinical staging.

#### **TESTIS PRESERVING SURGERY**

In patients with bilateral testicular germ cell tumors organ sparing surgery represents a new therapeutic approach with endocrinological and psychological advantages. Prerequisites need are the tumour should be organ confined with no infiltration of the rete testis, multiple biopsies of the tumour bed and peripheral parenchyma should be taken, any associated CIS should be treated by radiation therapy and patients must be followed closely (heidenreich – A et al 1997).

#### **LYMPHOMA**

Lymphoma is the most common secondary tumour of the testicle and the most frequent testicular neoplasm in men over the age of 50 years. Painless testicular enlargement is common, while bilateral involvement occurs in about third of patients.

Radical Orchiectomy establishes the diagnosis and cures a small subset of patients. Doxorubicin based regimen is used after orchiectomy. Survival is generally poor.

## **RESULTS**

All of our patients were treated surgically with high orchiectomy, and they had an uneventful stay in the hospital post operatively. All patients with Seminoma were treated by post operative Radiotherapy.

Chemotherapy with Cisplatin, Vincristine and Bleomycin was instituted for Teratoma and Embryonal cell carcinoma and both of these had 5 cycles each; following which they were lost to follow up.

Doxorubicin based chemotherapy Cisplatin, Adriamycin, Vincristine and cyclophosphamide was started for the 72 years old man with stage III non Hodgkins Lymphoma testis. This patient had one cycle of chemotherapy and is being followed up.

## CONCLUSION

On analyzing 324 cases of Scrotal swelling In GRH, Madurai over a period of 2 years the following characteristics were observed.

1. Primary vaginal hydrocele is the commonest cause for scrotal swellings and they commonly occur in the fourth decade
2. Secondary hydroceles account for a small percentage of the total and most of them were due to non specific epididymorchitis and usually at 30-40 years age group.
3. Post operative complications for hydrocele surgery were more after excision of sac than the other procedures; complications were very low for Lord's plication.
4. Varicoceles often tend to be left sided and their attribution to infertility is low in our series.
5. Non specific epididymoorchitis is the commonest cause for infection of the testis and epididymis.

6. Tuberculosis accounted for most of the cases of specific epididymo-orchitis.
7. Testicular tumors account for about 3.5% of all scrotal swelling and most of them were Semonomas (73%).
8. Detailed history and clinical examination alone were diagnostic in most of the cases, other investigations were all complimentary.
9. Testicular Ultrasonography in the diagnosis of Testicular tumor has a high sensitivity but the specificity is low.
10. Public should be made aware of the importance of self examination of the testis which might help in early diagnosis of Testicular tumor.

## **BIBLIOGRAPHY**

1. Ackermans Surgical Pathology Volume 2, 7<sup>th</sup> Edition: 949-988.
2. Anderson M et al ; tidsskr – Nor – Laegeforen 20: 113 (25) ; Oct 1993: 3146 – 7.
3. Anson Mcvay : Surgical Anatomy – 6<sup>th</sup> Edition 924 – 928
4. Aubert K et al “idiopathic Vaginal Hydrocele – Treatment using injection of Minicycline: Prog Uro 1993 april : 3(2) : 205 – 8.
5. Bailey & Love : Short practice of Surgery 22<sup>nd</sup> Edition : the testis and scrotum : Chapter 61 : 999-1010.
6. L.S. Baskin et al : Necrotising Soft Tissue Infection of the Perineum and Genitalia – BJU (1990), 65, 524-529.
7. Belloli et al : Laproscopic Surgery for Adolescent Varicocele : J. Paed Surgery 1996 Nov: 31 (11) : 1488 – 90.
8. J.P. Blandy : Testicular tumour – RA in Surgery by Selwyn taylor no 9, p. 24-268.
9. Calman F et al : B.J. Urology 51:154, 1979.
10. Charles M. Lindesy et al : Germinal Malignancies of the Testis : Journal of Urology Nov; Vol – 116, July 1976.
11. Chalouhy E et al : Microsurgical High Inguinal varicocelectomy with delivery of testis: J-Med-Liban 1994: 42:30:105-8.

12. Christopher L. Googen., M.D. et al : Bilateral testicular Tumour :Cancer 83/3.
13. A . Cuschieri : Essential Surgical Practice: 3<sup>rd</sup> edition 973, 1523-1528.
14. Dandappet et al : Effect of Hydrocele on testis and spermatogenesis : British Journal of Surgery Vol 77 : November : 1263-94.
15. Datta et al : Radionucleotide Imaging in Intra scrotal Lesions : J.A.M.A., 231 : 1975, 1960.
16. Donohue J.P: retro Lymphadenectomy : The antr Approach incl Bil. Suprarenal hilar Dissection : Urol. Clinic.North America 4:509, 1977.
17. Einhorn L.H.: Testicular Cancer as a Model for a Curative Neoplasm. The Richard and Hindal Rosenthal Foundation Award Lecture : Cancer Res 41:3275, 1981.
18. Einhorn L.H : Combination Chemotheraphy in Disseminated Testicular Cancer : the Indiana University Experience : Semin. Oncol, 6:82, 1979.
19. Elwyn E. Fraley : Testicular Tumour : the Urologic Clinics of NA : Vol 4 No 3: Oct 1977.
20. Farina LA et al : Treatment of Hydrocele with Evacuation and Percutaneous Sclerosis with Polydocanol : Actas-Urol-Esp. 194 June, 18\*6) : 690-3.
21. Farquharson's Text book of Operative Surgery : Eight Edition 672 – 683.

22. Fobbe F et al : Improvement in the Diagnosis of Scrotal Diseases using Colour Coded Duplex Sonography : Rofo-Fortcher-Gep-Ronygenster-Nukkamed : 1989 June : 150 (6) : 629-34.
23. Fuse H et al : Sclerotherapy of Hydroceles with Polydocanol : Int-Urol-Nephrol. 1994 : 26 (2) : 199-204.
24. George J. Bosl et al : Cancer of the testis : Cancer Principles and Practice of Oncology 5<sup>th</sup> Edition : Vincent T. Devita : 1397 – 1425.
25. Hahn et al : Testicular Scanning – A New Modality in Pre Operative Diagnosis of Testicular Torsion . J. Urol 125 : 213, 1981.
26. Handrikx A J et al : B mode Colour Duplex Sonography – A Useful Adjunct in Diagnosing Scrotal Disease : Br.J.Urol 1997 Jan 79 (1) 58-65.
27. Heidreich A et al : Testis Preserving Surgery in Bil. Testicular Germ Cell Tumours. Br.J.Urol. 1997 Feb : 79(2) : 253-7.
28. Holland et al : Conservative Management of Testicular Appendage Torsion. J. Urol 125 : 213, 1981.
29. Last's Anatomy Regional and Applied : 9<sup>th</sup> Edition 305 – 310.
30. Lee Mc Gregor's Synopsis of Surgical Anatomy : 12<sup>th</sup> Edition.
31. Lloyd M. Nyhus : Mastery of Surgery : 3<sup>rd</sup> Edition, 1658 – 1675.
32. Mostofi F.D : Testicular Tumour : Epidemiology, Aetiology & Pathologic Features : Cancer : 32: 1186 , 1976.



33. Musi. L et al : Modern trends in the Treatment of Varicocele : Paed. Med Chir 1998 Sep-Oct 18 (5suppl) : 31-34.
34. Oxford Text Book of Surgery : Vol.II 1576 – 1589, 1614-1617.
35. Oxford Text Book of Pathology : Vol II a 1544 – 1562.
36. Rados –N- et al : Biochemical aspects of Hydroceles : Acta. Med. Croatica 1996; 50 (1); 33-6.
37. Robbins, Kumar, Catran : Pathologic Basis of Disease 5<sup>th</sup> Edition : 1015 – 1022.
38. Rohandia O.P. et al : Everted Plication a Modified Surgery in Hydrocele : J. Postgrad. Med. 1993: Apr – June: 29 (2) : 77-8.
39. Russel.R.C.G : recent Advances in Surgery : 13.
40. Suwan P : Treatment of Hydcoceles by Aspiration & Tetracycline Instillation : J Med Asso Thai. 1994 August 77(8) : 421-5.
41. William C. Sharen M.D., : Acute Scrotal Pathologies : Surgical Clinics of North America Vol 62, Nov : 6 December 1982.
42. Campbell M.E. Walsh, P.C., Retik, A.B and Vaughan, E.D. (eds) (2002). Campbell's Urology 8<sup>th</sup> Edition
43. Sabistion Text book of Surgery (2004) 17<sup>th</sup> Edition.
44. Schwartz : Principles of Surgery 7<sup>th</sup> Edition.

**PROFORMA**  
***DISSERTATION ON SCROTAL SWELLINGS***  
***(Undescended Testis and Hernia Excluded)***

|            |   |           |
|------------|---|-----------|
| Name       | : | I.P. No.: |
| Age        | : | D.O.A. :  |
| Occupation | : | D.O.S. :  |
| Address    | : | D.O.D. :  |
| Duration   | : |           |

Onset      Acute  
              Subacute  
              Chronic

H/O    1. Trauma  
         2. Fever with Chills  
         3. Cough with expectoration / COPD /PT  
         4. Diabetes  
         5. Associated Medical Diseases if any  
         6. Pain                      Mild / Moderate / Severe  
         7. Exposure History  
         8. Family History  
         9. Any other symptoms

**GENERAL EXAMINATION**

Febrile / Anaemic / Edema feet / Clubbing / Gynecomastic / Jaundice /

Lymph node

Pulse

BP

**LOCAL EXAMINATION**                      - Scrotal Swelling

|      |                                 |
|------|---------------------------------|
| Side | - Size                          |
| Skin | - Normal / thickened / rugosity |

Signs of inflammation

Scrotal edema

Sinus / Ulcer / Redness / Blackening

Foul smell

Upper limit of the swelling

Reducible / Tenderness / Warmth

Skin – prefixed

Sinus - Fixed or not

Number

Anterior / Posterior

Discharge

Testis - Swollen

Normal

Atrophied

Sensation

Lie

If there is swelling

- |                           |   |                                   |
|---------------------------|---|-----------------------------------|
| 1. Consistency            | - | Cystic / Solid                    |
| 2. Fluctuation            | - | +ve / -ve                         |
| 3. Transillumination      | - | +ve / -ve                         |
| 4. Cough impulse          | - | +ve / -ve                         |
| 5. Tenderness             | - | +ve / -ve                         |
| 6. Epididymis             | - | Swollen / Nodules / Number        |
| 7. Vas deferens – Nodules | - | Present / Absent                  |
| 8. Cremaster              | - | Hypertrophied / Not hypertrophied |
| 9. Any other appearance   | - | Bag of worms                      |
| 10. RLNS                  | - | Inguinal / Iliac / Paraaortic     |

Abdomen – any mass – paraortic nodes

Liver          Spleen

Kidneys      Ascites

P / R

RS – Effusion / Fibrosis / Collapse / Normal

CVS –

Virchows node – Palpable / Not

## INVESTIGATIONS

Hb%                      TC                      DC                      ESR

Peripheral smear for Mf

Blood Urea

Blood Sugar

VDRL

HIV

Microscopic examination of discharge

Culture

X-ray of Scrotum

X-ray Chest

US Scan

Conservative treatment – specify the drugs and other measures

Pre Op findings

    Anaesthesia

    Incision

Sac – Normal / Thickened / Calcified

Content              - Clear fluid

                         Altered blood

Blubbery material

Pus

Chyle

Smell

Testis - Normal / Atrophied / Pathological / Colour

If tumor - Cut section

Epididymis - Normal / Nodular

Vas deferens - Normal / Beading if any

Nature of surgery

- Eversion of sac
- Excision of sac
- Plication of sac
- Low Orchiectomy
- High Orchiectomy

Follow up